FOR VETERINARY USE ONLY

ROMDOLL

SHEEP-GOAT POX VACCINE

Dollvet

LYOPHILISATE SUSPENSION FOR INJECTION AND SOLVENT

COMPOSITION:

Following minimum quantities are used in each dose (1ml).

Ingredients	Quantity	Strength
Attenuated RM65 strain	$10^{2,5}$ $TCID_{50}$/dose	Immunizing antigen
Lactalbumin hydrolysate	0,25 mg/dose	Preservative
Sucrose	0,50 mg/dose	

INDICATIONS AND CONTRAINDICATIONS:

ROMDOLL is used against sheep-goat pox, that causes severe generalized infections in sheep, goats, lambs and kids, and immunity occurs in target animals through a cellular and humoral immune response 21 days after vaccination. It has no contraindications.

ADMINISTRATION AND DOSAGE:

The vaccine is administered by subcutaneous injection into the axillary area in the dose of 1 ml for sheep and goats and 0,5 ml for lambs and kids.

WARNINGS:

Using sterile syringes, the lyophilized pellet should be homogenized, thoroughly dissolving it with the reconstitution solvent. During vaccination, special care should be taken not to contaminate the eyes or hands. In case of an accidental contact, the site should be cleaned with water and disinfected.

ADVERSE REACTIONS:

In some animals, hard or soft swellings of 0,2-5 cm in diameter may develop at the vaccination site, and a temporary rise in body temperature may be observed.

WITHDRAWAL PERIODS:

0 day.

STORAGE CONDITIONS:

Real time shelf life is 2 years at + 2/8ºC and in the dark.

PRESENTATION:

For vaccine lyophilization; 1-2 ml of 50, 100 and 200 doses of vaccine suspension will be placed in Type I neutral glass bottles of 5 and 10 ml in volume, and butyl stoppers and aluminum caps will be used to close the bottles. For reconstitution solvent; 50 ml, 100 ml and 200 ml Type II neutral glass or plastic bottles. Butyl stoppers and aluminum caps are used to close the bottles.

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DISPOSAL PRECAUTIONS:

Unused or opened but not completely used vaccines are disposed of as medical waste, according to the provisions of the applicable waste regulations..

"Keep out of reach of children" and "Consult a veterinarian upon an unexpected effect"

LEAFLET APPROVAL DATE: 17.05.2017

DATE OF MARKETING AUTHORIZATION ISSUED BY MINISTRY OF AGRICULTURE AND FORESTRY: 17.05.2017

NAME AND ADDRESS OF MARKETING AUTHORIZATION HOLDER:

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NAME AND ADDRESS OF PRODUCTION SITE:

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